## Commonwealth of Virginia Department of General Services Division of Consolidated Laboratory Services Richmond, Virginia

## Method Detection Limit Revision 2 (40 CFR 136 App B)

Definition and Procedure for the Determination of the Method Detection Limit, Revision 2 [40 CFR 136 Appendix B] EPA 821-R-16-006, https://www.epa.gov/cwa-methods						
The method detection limit (MDL) is defined as the minimum measured concentration of a substance that can be reported with 99% confidence that the measured concentration is distinguishable from method blank results.						
Facility Name:VELAP ID						
Assessor Name:Analyst Name:	Inspe	ectio	n Da	ate	·····	
SUMMARY CHECKLIST (see full checklist for supporting details and full compliance):  All sample processing steps Initial: 7 samples + 7 blanks minimum Prep/Testing on at least 3 days Testing uses all instruments Determined MDL is greater of MDL <sub>S</sub> and MDL <sub>B</sub> Ongoing: 7 samples (same spike conc) + 7 blanks minimum over year Collected each quarter samples were run (2 determinations per instrument, separate batches) Spiking level increased (and initial repeated) if >5% of spikes fail to return positive numerical results / meet all identification criteria Uses all data for same level spike and all blank data (or can use last 6 months or 50 most recent, whatever is greater) Annual re-calculation (<13 months) and evaluation of MDL <sub>S</sub> and MDL <sub>B</sub> using most recent 24 months' data Verified correct Student's T value used MDL verified as within 0.5 to 2.0 times the existing MDL with <3% of blanks above existing MDL, or initial is repeated For simultaneous 2016 TNI Standard Compliance (V1M4 1.5.2): use spike level at or below LOQ Supporting data available as specified						
Relevant Aspect of Standards	Method Reference	Y	N	N/A	Comments	
Records Examined: SOP Number/ Revision/ Date			,	Analyst	:	
Method: Matrix:	Analy	/te: _			· · · · · · · · · · · · · · · · · · ·	
Instrument(s):	(s): MDL Date:					
Were all sample processing steps used by the laboratory included in the determination of the MDL?	Scope and Application					
2. NOTE: The MDL is not applicable to methods where low-level spiked samples cannot be prepared. An MDL may be based on method blanks for gravimetric methods. (Refer to method for additional detail.)  An MDL will not be expected by VELAP for: BOD/CBOD, color, odor, pH, alkalinity, specific conductance, turbidity, total solids or residue, total (TS), total suspended solids or residue, non-filterable (TSS),	Scope and Application					

Document #:21799 Revision: 5

			1	T
e, volatile (TVS), dissolved				
	Scope and Application			
NITIAL MDL				
the selection(s)) termined concentration plus 3x of a set of method blanks ation value that corresponds to noise ratio in the range of 3 to 5 ation equivalent to three times the eplicate instrumental ed blanks of the calibration where there is a ensitivity, i.e, a break in the slope imitations	1(a) through 1(f)			
HE INITIAL MDL				
e data to perform the Ongoing cified in Section (4), typically implemented or if a method was	2			
Is in excess of 10x the estimated e required for analytes with very ning at this stage so that the neets the requirements of 2016 II enable the laboratory to use the	2(a)			
spiked samples and 7 method	2(b)			
	2(b)			
	2(b)			
pe used, if compliant with the at least 3 batches and generated months.	2(b), 2(b)(iii)			
	or residue, filterable (TDS), total e, volatile (TVS), dissolved ure.  Id from a clean reference matrix and consistent quantity of the  NITIAL MDL  Bestimated using one or more of the selection(s)) termined concentration plus 3x of a set of method blanks ation value that corresponds to noise ratio in the range of 3 to 5 ation equivalent to three times the eplicate instrumental and blanks of the calibration where there is a tensitivity, i.e, a break in the slope dimitations termined MDL  THE INITIAL MDL  Lis used when the laboratory e data to perform the Ongoing ciffed in Section (4), typically implemented or if a method was at months  2-10 times the estimated MDL  Is in excess of 10x the estimated are required for analytes with very oning at this stage so that the meets the requirements of 2016 are required for analytes with very oning at this stage so that the meets the requirements of 2016 are to satisfy 2016 TNI V1M4 1.5.2. Spiked samples and 7 method and all steps of the method?  The MDL prepared in at least 3 at a calendar dates?  The MDL analyzed on three is a least 3 batches and generated a months.  The MDL analyzed on three is a least 3 batches and generated a months.  The MDL analyzed on three is a least 3 batches and generated a months.  The MDL analyzed on three is a least 3 batches and generated a months.  The MDL analyzed on three is a least 3 batches and generated a months.  The MDL analyzed on three is a least 3 batches and generated a months.  The MDL analyzed on three is a least 3 batches and generated a months.  The MDL analyzed on three is a least 3 batches and generated a months.	d from a clean reference matrix and consistent quantity of the  NITIAL MDL  estimated using one or more of the selection(s)) termined concentration plus 3x of a set of method blanks at ornesponds to enoise ratio in the range of 3 to 5 ation equivalent to three times the explicate instrumental and blanks for the calibration where there is a ensitivity, i.e., a break in the slope dimitations termined MDL  THE INITIAL MDL  is used when the laboratory end ata to perform the Ongoing cified in Section (4), typically implemented or if a method was at months  2-10 times the estimated MDL  Is in excess of 10x the estimated the required for analytes with very analysis and 7 method and all steps of the method?  The MDL prepared in at least 3 are calendar dates?  The MDL analyzed on three size and so the same day, be used, if compliant with the at least 3 batches and generated a months.  Scope and Application  Italian plus 3 to 5 at 5	d from a clean reference matrix and consistent quantity of the settimated using one or more of he selection(s)) termined concentration plus 3x of a set of method blanks ation value that corresponds to noise ratio in the range of 3 to 5 ation equivalent to three times the applicate instrumental ad blanks if the calibration where there is a ensitivity, i.e., a break in the slope limitations itermined MDL  THE INITIAL MDL  Is is used when the laboratory e data to perform the Ongoing cified in Section (4), typically implemented or if a method was 24 months  2-10 times the estimated MDL  Is in excess of 10x the estimated he required for analytes with very  2(a)  ning at this stage so that the neets the requirements of 2016 III enable the laboratory to use the to satisfy 2016 TNI V1M4 1.5.2. spiked samples and 7 method gh all steps of the method?  If the MDL prepared in at least 3 ate calendar dates?  If the MDL analyzed on three is?  2(b)  analysis may be on the same day, be used, if compliant with the rate least 3 batches and generated at months.	e, volatile (TVS), dissolved ure.  d from a clean reference matrix and consistent quantity of the  NITIAL MDL  estimated using one or more of the selection(s)) termined concentration plus 3x of a set of method blanks attion value that corresponds to enoise ratio in the range of 3 to 5 attion equivalent to three times the applicate instrumental ad blanks for the calibration where there is a ensitivity, i.e, a break in the slope limitations termined MDL  THE INITIAL MDL  Is is used when the laboratory edata to perform the Ongoing cified in Section (4), typically implemented or if a method was 24 months  2-10 times the estimated MDL is in excess of 10x the estimated her required for analytes with very  also analytes with very  2(a)  ning at this stage so that the neets the requirements of 2016 ill enable the laboratory to use the to satisfy 2016 TNI V1M4 1.5.2. spiked samples and 7 method righ all steps of the method?  The MDL prepared in at least 3 ate calendar dates?  2(b)  ate calendar dates?  2(b)  also small stage in the same day be used, if compliant with the at at least 3 batches and generated at months.

and spiked samples must be used. Only data associated with gross failures with documentation for rationale (ex: instrument malfunctions, mislabeled samples, cracked vials) may be removed.  o The same prepared extract may be analyzed on multiple instruments.  o A spiked sample and a method blank sample may be analyzed in the same batch, but are not required to be.	
11. If multiple instruments will be assigned the same MDL, sample analyses must be distributed across all instruments. Was each instrument represented with a minimum of two spiked samples and two blank samples prepared / analyzed on different days?	2(b)(i), 2(b)(ii), 2(d)
12. Spiking level evaluation: Did every result from spiked samples meet the method qualitative identification criteria? Did every result from spiked samples provide a numerical result greater than zero?  If the answer to either question is NO, the spiked samples used for initial MDL determination must be repeated at a higher concentration.	2(c)
13. Were all computations made as specified in the analytical method and expressed in the method-specified reporting units?	2(d)
14. Was the MDL <sub>s</sub> (the MDL based on spiked samples) computed as follows?	2(d)(ii)
15. Was the MDL <sub>b</sub> (the MDL based on method blanks) computed as follows?  If none of the method blanks give numerical results for an individual analyte, the MDL <sub>b</sub> does not apply.  NOTE: A numerical result includes both positive and negative results, including results below the current MDL, but not results of "ND" [not detected] commonly observed when a peak is not present in chromatographic analysis.  OR	2(d)(iii)(A)
If some (but not all) of the method blanks for an individual analyte give numerical results, set the MDL <sub>b</sub> equal to the highest method blank result.	2(d)(iii)(B)

If more than 100 method blanks are available, set MDL <sub>b</sub> to the level that is no less than the $99^{th}$ percentile of the method blank results. For "n" method blanks where $n \ge 100$ , sort the method blanks in rank order. The $(n * 0.99)$ ranked method blank result (round to the nearest whole number) is the MDL <sub>b</sub> . [Refer to published method for a mathematical example.] OR  If all of the method blanks for an individual anayte give numerical results, then calculate the MDL <sub>b</sub> as: $ MDL_b = \overline{x} + t_{(n-1, 1-\alpha=0.99)} S_b $ Where $ MDL_b = MDL  based on method blanks $ $ \overline{x} = the mean of the method blank results $ $ t_{(n-1, 1-\alpha=0.99)} = the Student's t-value appropriate for a single-tailed 99^{th} percentile t statistic and a standard deviation estimate with n-1 degrees of freedom. (See Table 1, below; 3.143 when n=7) Sb = sample standard deviation of the replicate method blank analyses  NOTE: If the mean of the blanks is <0 (i.e., a negative number), substitute 0 for the mean.  NOTE: If 100 or more method blanks are available, as an option, MDL_b may be set to the concentration that is greater than or equal to the 99^{th} percentile of the method blank results, as described in Section (2)(d)(iii)(B)$	2(d)(iii)(C)
16. Was the greater of MDL <sub>s</sub> or MDL <sub>b</sub> selected as the <u>initial MDL</u> ?	2(e)
ONGOING DATA COLLECTION	
17. Was ongoing data collected as follows?	
During any quarter in which samples are being analyzed, prepare and analyze a minimum of two spiked samples on each instrument, in separate batches, using the same spiking concentration used in Section 2 [initial MDL calculation].  NOTE: If any analytes are repeatedly not detected in the quarterly spiked sample analyses or do not meet the qualitative identification criteria of the method the spiking level should be adjusted upward. (See 3(c).)  NOTE: It is not necessary to analyze additional method blanks together with spiked samples; include all of the routine method blanks analyzed with each	3(a)

batch during the course of (routine) sample analysis.	
18. Did ongoing data collection ensure that at least seven spiked samples and seven method blanks were completed for the annual verification?  NOTE: If only one instrument is in use, a minimum of seven spikes are still required, but they may be drawn from the last two years of data collection.	3(b)
<ul> <li>19. At least once per year, was the spiking level reevaluated?</li> <li>NOTE: If more than 5% of the spiked samples do not return positive numerical results that meet all method qualitative identification criteria, the spiking level must be increased and the initial MDL re-determined following the procedure in Section 2.</li> </ul>	3(c)
20. NOTE: If the method is altered in a way that can be reasonably expected to change its sensitivity, redetermine the initial MDL according to Section 2 and restart the ongoing data collection.	3(d)
<ul> <li>21. If applicable, was the following addressed if a new instrument was added?  If a new instrument is added to a group of instruments whose data are being pooled to create a single MDL, analyze a minimum of two spiked replicates and two method blank replicates on the new instrument.  • If both method blank results are below the existing MDL, then the existing MDL<sub>b</sub> is validated.  • Combine the new spiked sample results to the existing spiked sample results and recalculate the MDL, as in Section 4.</li> <li>• If the recalculated MDL<sub>s</sub> does not vary more than the factor specified in Section 4(f) of this procedure, then the existing MDLs is validated.</li> <li>• If either of these two conditions is not met, then calculate a new MDL following instructions in Section 2.</li> </ul>	3(e)
ONGOING ANNUAL VERIFICATION	
22. Was the MDL <sub>s</sub> and MDL <sub>b</sub> re-calculated at least once every thirteen months from the collected spiked samples and method blank results from the last 24 months using the equations in Section 2?	4(a)
23. For the MDL <sub>s</sub> , was all data generated within the last 24 months, but only data with the same spiking level, included in the recalculation?	4(b), 4(c), 4(d)

	NOTE: Include the initial MDL spiked samples, if the data were generated within 24 months.			
	NOTE: Use only data associated with acceptable calibrations and batch QC. Include all routine data with the exception of batches that are rejected and the associated samples reanalyzed.			
	NOTE: Only documented instances of gross failures may be excluded from the calculations.			
	NOTE: If the laboratory believes the sensitivity of the method has changed significantly, then the most recent data available (i.e., data collected after the change) may be used, maintaining compliance with the requirement for at least 7 replicates in three batches on three separate days (per Section 2(b).)			
24.	For the MDL <sub>b</sub> , were all method blank results from the last 24 months used?			
	NOTE: The laboratory has the option to use only the last six months of method blank data or the 50 most recent method blanks, whichever criteria yields the greater number of method blanks.	4(e)		
	Indicate the option used by the laboratory:			
25.	Was the verified MDL the greater of the MDL $_{\! s}$ or MDL $_{\! b}?$	4(f)		
26.	Was the verified MDL within 0.5 to 2.0 times the existing MDL, and did fewer than 3% of the method blank results have numerical results above the existing MDL?			
	If so, the existing MDL may be left unchanged at the option of the laboratory.	4(f)		
	If not, adjust the MDL to the new verified MDL.			
27.	NOTE: Refer to the published method for determination of the MDL for a specific (sample) matrix.	Addendum		
28.	Were documentation requirements met?  The prep date, analysis date, and instrument for each analysis was available for evaluation of MDL compliance.  The analytical method used for MDL determination was specifically identified by number or title.  The MDL for each analyte was expressed in the method reporting units.  Data and calculations used to establish the MDL	Documentation and Procedure		
	can be reconstructed upon request The sample matrix used to determine the MDL			

was identified with the MDL value The mean spiked and recovered analyte levels were documented with the MDL The rationale for removal of outlier results, if any, was documented and maintained on file with the results of the MDL determination.		
Notes/ Comments:		
Notes/ Comments:		

THIS SCHECKLIST IS AN INTERVIEW AND/OR DATA REVIEW TOOL USED BY ASSESSORS AND IS NOT TO BE CONSIDERED AS A SUBSTITUTE FOR REQUIREMENTS OF THE PUBLISHED METHOD. CHECKLISTS ARE SUBJECT TO CHANGE. PLEASE NOTIFY DCLS IMMEDIATELY BY EMAIL OF ANY IDENTIFIED ERRORS OR OMISSIONS. (Lab Cert@dgs.virginia.gov)

Table 1: Single-Tailed 99<sup>th</sup> Percentile *t* Statistic

Number of replicates	Degrees of freedom (n-1)	t <sub>(n-1, 0.99)</sub>
7	6	3.143
8	7	2.998
9	8	2.896
10	9	2.821
16	15	2.602
32	31	2.453
50	49	2.405
80	79	2.374
100	99	2.365

A Student's T table with values to 100 and a calculation tool for values >100 is located on the VELAP toolbox webpage at <a href="https://www.dgs.virginia.gov/dcls">www.dgs.virginia.gov/dcls</a> (Choose Laboratory Accreditation, then Toolbox), or use this <a href="https://www.dgs.virginia.gov/dcls">LINK</a>.

Title: Method Detection Limit Revision 2 (40 CFR 136 App B)

Document #:21799

Revision: 5